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ASSISTANT COMMISSIONER FOR PATENTS
Washington, D.C. 20231

Sir:

Transmitted herewith for filing under 37 C.F.R. §1.53(b) is the patent application of:
Inventor(s): Gertrud HÖTTEN, Helge NEIDHARDT, and Jen POHL

For: **NEW GROWTH/DIFFERENTIAL FACTOR OF THE TGF- β FAMILY**

This application is a divisional of Application No. 08/288,508

☒ Return Receipt Postcard

☒ Specification (37 pages)

☒ 7 sheets of drawings

☒ A Sequence Listing and Statement with disk

☒ A Preliminary Amendment

☒ Declaration and Power of Attorney

☒ Copy from a prior application

☒ The disclosure of the prior application, from which a copy of the declaration is supplied as noted above is considered as being a part of the disclosure of the accompanying application and is hereby incorporated by reference therein.

☒ Assignment was filed in parent Application No. 08/288,508, Reel/Frame 7252/0028.

☒ Priority of German applications, Serial Nos. P43 26 829.3 filed 08/10/1993, P44 18 222.8 filed 05/25/1994, and P44 20 157.5 filed 06/09/1994 is claimed under 35 U.S.C. §119.

☒ An Information Disclosure Statement with PTO-1449

☒ A filing fee, calculated as shown below, including claims added or deleted in the Preliminary Amendment (Check No. 20646):

(Col. 1)			(Col. 2)		Small Entity		Other Than A Small Entity	
FOR:	No. Filed	No. Extra	RATE	FEE		RATE	FEE	
BASIC FEE				\$380	or		\$760.	
TOTAL CLAIMS	12 - 20 =	-0-	× 9 =		or	× 18 =	-0-	
INDEP CLAIMS	4 - 3 =	1	× 39 =		or	× 78 =	78.	
_ MULTIPLE DEPENDENT CLAIM PRESENTED			+130 =		or	+260 =	-0-	
			TOTAL		or		\$848.	

Respectfully submitted,
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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:

HÖTTEN et al

Art Unit: Unassigned

Serial Number: Unassigned

Examiner: Unassigned

Filed: August 31, 1999

For: NEW GROWTH/DIFFERENTIATION FACTOR OF THE TGF- β FAMILY

PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, D.C. 20231

August 31, 1999

Sir:

Prior to calculation of the filing fee and prior to the examination of this application, please amend the above-identified application as follows:

IN THE SPECIFICATION

Please insert new pages 36-52 containing the revised Sequence Listing and renumber the pages containing the claims and abstract accordingly.

IN THE CLAIMS:

Kindly cancel claim 1 without prejudice or disclaimer.

Please add the following new claims to the application:

--13. A protein of the TGF- β family encoded by an isolated DNA molecule which has a sequence selected from the group consisting of:

- a) the sequence shown in SEQ ID NO:1,
- b) a part of SEQ ID NO:1 which encodes the mature protein,

c) a nucleotide sequence which encodes the amino acid sequence according to SEQ ID NO:2,

d) a nucleotide sequence which encodes a portion of the amino acid sequence according to SEQ ID NO:2, wherein said portion is the mature protein, and

e) a sequence which hybridizes to a), b), c) or d) under stringent hybridization conditions, wherein said sequence encodes the mature protein of the TGF- β family.

14. The protein according to claim 13, wherein said protein has the amino acid sequence shown in SEQ ID NO:2.

15. A pharmaceutical composition comprising at least one protein according to claim 13, in combination with a pharmaceutically acceptable carrier, auxiliary substance, diluent or filler.

16. A method for treating or preventing damage to bone, cartilage, connective tissues, skin, mucous membranes, epithelium or teeth, comprising administering the protein according to claim 13 to a patient in need of such treatment.

17. A method for improving wound healing and tissue regeneration, comprising administering the protein according to claim 13 to a patient in need of such treatment.

18. A dental implant which contains the protein according to claim 13.

19. An antibody or antibody fragment which binds to the protein according to claim 13.

20. An antibody or antibody fragment which binds to a protein comprising the amino acid sequence shown in SEQ ID NO:2 or biologically functional parts thereof.

21. A pharmaceutical composition according to claim 15, wherein the protein is contained on or in a natural or synthetically prepared matrix material.

22. A pharmaceutical composition according to claim 21, wherein said matrix material is a biocompatible porous material that can be biologically degraded.

23. A protein of the TGF- β family encoded by a DNA molecule which is a part of SEQ ID NO:1 which encodes the mature protein and further comprises nucleotide sequences coding for signal and/or propeptide parts.

24. A protein of the TGF- β family encoded by a DNA molecule which comprises nucleotides 640-2142 of SEQ ID NO:1.--

REMARKS

The above amendments have been made to put the application into better condition for examination.

In this amendment, applicants also amend the specification by inserting a Sequence Listing as pages 36-52 and renumbering the pages containing the claims and abstract accordingly.

It is respectfully submitted that the foregoing amendments and remarks place the application in compliance with 37 CFR 1.821 through 1.825 and no new matter is contained in this amendment.

Enclosed is a computer-readable sequence listing in MS-DOS format on a 3.5" disk. The filename is 5649022.APP.

Should any additional information be required, please contact applicants' undersigned representative at the telephone number listed below.

In the event that any fees are due in connection with this paper, please charge our Deposit Account No. 14-1060.

In the event this paper is not being timely filed, the Applicants respectfully petition for an appropriate extension of time. Any fees for such an extension together with any additional fees may be charged to Counsel's Deposit Account 14-1060.

Respectfully submitted,

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New growth/differentiation factor
of the TGF- β family

DESCRIPTION

The present invention concerns a new growth/differentiation factor of the TGF- β family and DNA sequences coding therefor.

The TGF- β family of growth factors which includes BMP-, TGF- and inhibin-related proteins (Roberts and Sporn, Handbook of Experimental Pharmacology 95 (1990), 419-472) is particularly relevant for a wide range of medical treatment methods and applications. These factors are suitable in methods which concern wound-healing and tissue regeneration. Furthermore several members from the TGF- β family induce tissue growth, in particular growth of bones, and therefore play a crucial role in inducing the development of cartilage and bones.

Wozney (Progress in Growth Factor Research 1 (1989), 267-280) and Vale et al (Handbook of Experimental Pharmacology 95 (1990), 211-248) describe various growth factors such as those which are related to the BMP group (bone morphogenetic proteins) and the inhibin group. The members of these groups show significant structural similarities. The precursor of the protein consists of an amino-terminal signal sequence, a propeptide and a carboxy-terminal sequence of about 110 amino acids that are cleaved from the precursor and constitute the mature protein. In addition their members are defined by an amino acid sequence homology. The mature protein contains the most conserved sequences, in particular seven cysteine residues which are conserved among the family members. The TGF- β -like proteins are multifunctional, hormonally active growth factors. They also have related biological activities such as

chemotactic attraction of cells, promotion of cell differentiation and tissue-inducing capabilities such as cartilage-inducing and bone-inducing capabilities. The US Patent No. 5,013,649 discloses DNA sequences that code for osteo-inductive proteins that are denoted BMP-2 and the US Patent Applications series No. 179 101 and 170 197 disclose the BMP proteins BMP-1 and BMP-3. Moreover many types of cells are able to synthesize TGF- β -like proteins and practically all cells have TGF- β receptors.

On the whole these proteins show differences in their structure which leads to significant variations in their exact biological function. In addition they are found in a wide range of different types of tissue and at various stages of development. As a result they can exhibit differences with regard to their exact function e.g. the required cellular physiological environment, their life-span, their target sites, their requirements for auxiliary factors and their stability against degradation. Thus, although a multitude of proteins have been described that exhibit a tissue-inductive and in particular osteo-inductive potential, their natural functions in the organism and - more significantly - their medical relevance still have to be investigated in detail. It is thought to be highly probable that members of the TGF- β family are present that are still unknown which are important for osteogenesis or the differentiation/induction of other types of tissue. A major difficulty in the isolation of these new TGF- β -like proteins is, however, that their functions cannot yet be described exactly enough to develop highly discriminating bioassays. On the other hand the expected nucleotide sequence homology to other members of the family is too low to enable a screening by classical

nucleic acid hybridization techniques. Nevertheless the further isolation and characterization of new TGF- β -like proteins is urgently required in order to provide further inducing and differentiation-promoting proteins that fulfil all the desired medical requirements. These factors could be used medically in the healing of lesions and the treatment of degenerative diseases of bones and/or other types of tissue such as the kidney or the liver.

A nucleotide and amino acid sequence for the TGF- β protein MP-52 is stated in the Patent Application PCT/EP93/00350 in which the sequence corresponding to the mature peptide and a major portion of the sequence corresponding to the propeptide of MP-52 are given. The complete sequence of the propeptide MP-52 is not disclosed.

The object on which the present invention is based is to provide DNA sequences that code for new members of the TGF- β protein family with mitogenic and/or differentiation-inductive e.g. osteo-inductive potential. The object of the present invention was therefore in particular to provide the complete DNA and amino acid sequence of the TGF protein MP-52.

This object is achieved by a DNA molecule that codes for a protein of the TGF- β family and which comprises

- (a) the part coding for the mature protein and if desired further functional parts of the nucleotide sequence shown in SEQ ID NO. 1,
- (b) a nucleotide sequence corresponding to a sequence from (a) within the scope of the degeneracy of the genetic code,

(c) an allelic derivative of a nucleotide sequence corresponding to one of the sequences from (a) and (b) or

(d) a sequence hybridizing with one of the sequences from (a), (b) or (c) provided that a DNA molecule according to (d) contains at least the part coding for a mature protein from the TGF- β family.

Further embodiments of the present invention concern the subject matter of claims 2 to 10. Other features and advantages of the invention can be derived from the description of the preferred embodiments and figures. The sequence protocols and figures are now briefly described.

SEQ ID NO. 1 shows the complete nucleotide sequence of the DNA coding for the TGF- β protein MP-52. The ATG start codon starts with nucleotide 640. The start of the mature protein begins after nucleotide 1782.

SEQ ID NO. 2 shows the complete amino acid sequence of the TGF- β protein MP-52 which was derived from the nucleotide sequence shown in SEQ ID NO. 1.

Figure 1 shows a comparison between the amino acid sequence of MP-52 and several members of the BMP protein family starting with the first of the seven conserved cysteine residues. * denotes that the amino acid is the same in all compared proteins; + denotes that the amino acid corresponds in at least one of the proteins compared to MP-52.

Figure 2 shows the nucleotide sequences of the oligonucleotide primers that were used in the present invention and a comparison of these sequences with known members of the TGF- β family. M denotes A or C, S denotes C or G, R denotes A or G and K denotes G or T. 2a shows the sequence of the primer OD, 2b shows the sequence of the primer OID.

The present invention encompasses at least the part coding for the mature protein and if desired further functional parts of the nucleotide sequence shown in SEQ ID NO.1 as well as sequences that correspond to this sequence within the scope of the degeneracy of the genetic code and allelic derivatives of such sequences. In addition the present invention also encompasses sequences that hybridize with such sequences provided that such a DNA molecule completely contains at least the part coding for the mature protein of the TGF- β family.

The term "functional part" within the sense of the present invention denotes a protein part which is capable of acting for example as a signal peptide, propeptide or as a mature protein part i.e. it fulfils at least one of the biological functions of the natural protein parts of MP-52.

The region coding for the mature part of the protein extends from nucleotides 1783 - 2142 of the sequence shown in SEQ ID NO.1. If desired, the DNA molecule can also comprise further functional parts of the sequence shown in SEQ ID NO. 1, namely the nucleotide sequences coding for the signal or/and propeptide part. It is particularly preferred that the DNA molecule comprises

the sequence for the signal and the propeptide part and the part of the mature protein i.e. nucleotides 640-2142 of the sequence shown in SEQ ID NO. 1. On the other hand the DNA molecule can also comprise functional signal or/and propeptide parts from other proteins in addition to the part coding for the mature protein, in particular from other proteins of the TGF- β family e.g. the above-mentioned BMP proteins. The respective nucleotide sequences are given in the references mentioned above to the disclosure of which reference is hereby being made.

Moreover the present invention also encompasses a DNA molecule as defined above that contains a non-coding intron sequence between nucleotides 1270 and 1271 of the sequence shown in SEQ ID NO. 1. This intron sequence is contained in the plasmid SKL 52 (H3) MP12 which is deposited at the DSM and has the genomic nucleic acid sequence of MP-52.

The invention also encompasses the cDNA sequence of the MP-52 protein coded by the phage λ 15.1. This sequence starts with nucleotide 321 of SEQ ID NO. 1.

Although the allelic, degenerate and hybridizing sequences which are encompassed by the present invention have structural differences due to slight changes in their nucleotide or/and amino acid sequence, the proteins coded by such sequences still essentially have the same useful properties that enable their use in basically the same medical applications.

The term "hybridization" according to the present invention means the usual hybridization conditions, preferably conditions with a salt concentration of 6 x

SSC at 62 to 66°C followed by a one hour wash with 0.6 x SSC, 0.1 % SDS at 62 to 66°C. It is particularly preferred that the term "hybridization" denotes stringent hybridization conditions with a salt concentration of 4 x SSC at 62 to 66°C followed by a one hour wash with 0.1 x SSC, 0.1 % SDS at 62 to 66°C.

Preferred embodiments of the present invention are DNA sequences as defined above that are obtainable from vertebrates, preferably mammals such as pigs, cows and rodents such as rats or mice and in particular from primates such as humans.

A particularly preferred embodiment of the present invention is the sequence denoted MP-52 shown in SEQ ID NO. 1. The transcripts of MP-52 were obtained from embryonic tissue and code for a protein which has a considerable amino acid homology to the mature portion of BMP-like proteins (see Fig. 1). The protein sequences of BMP2 (=BMP2A) and BMP4 (=BMP2B) are described by Wozney et al., Science 242 (1988), 1528-1534. The corresponding sequences of BMP5, BMP6 and BMP7 are described by Celeste et al., Proc. Natl. Acad. Sci. USA 87 (1990), 9843-9847. Several typical sequence homologies which are specific for known BMP sequences have also been found in the propeptide part of MP-52 whereas other parts of the precursor part of MP-52 exhibit considerable differences to BMP precursors.

In addition the present invention concerns a vector that contains at least one copy of a DNA molecule according to the invention. The DNA sequence according to the invention is preferably operatively linked to an expression control sequence in such a vector. Such

vectors are suitable for the production of TGF- β -like proteins in stably or transiently transformed cells. Various animal, plant, fungal and bacterial systems can be used for the transformation and subsequent culture. The vectors according to the invention preferably contain sequences necessary for replication in the host cell and are autonomously replicable. Furthermore the use of vectors is preferred that contain selectable marker genes which can be used to detect transformation of a host cell.

In addition the invention concerns a host cell which is transformed with a DNA according to the invention or with a vector according to the invention. Examples of suitable host cells include various eukaryotic and prokaryotic cells such as *E. coli*, insect cells, plant cells, mammalian cells and fungi such as yeast.

In addition the invention concerns a protein of the TGF- β family which is coded by a DNA sequence according to claim 1. The protein according to the invention preferably has the amino acid sequence shown in SEQ ID NO. 2 or if desired functional parts thereof and exhibits biological properties such as tissue-inductive in particular osteo-inductive or/and mitogenic capabilities that may be relevant for a therapeutic application. The above-mentioned characteristics of the protein can vary depending on the formation of homodimers or heterodimers. Such structures may also prove to be suitable for clinical applications.

The biological properties of the proteins according to the invention, in particular the mitogenic and osteo-inductive potential can be determined for example in

assays according to Seyedin et al., PNAS 82 (1985), 2267-2271 or Sampath and Reddi, PNAS 78 (1981), 7599-7603.

Furthermore the present invention concerns a process for the production of a protein of the TGF- β family which is characterized in that a host cell transformed with a DNA according to the invention or with a vector according to the invention is cultured and the TGF- β protein is isolated from the cell or/and culture supernatant. Such a process comprises culturing the transformed host cell in a suitable culture medium and purifying the TGF- β -like protein produced. In this manner the process enables the production of an adequate amount of the desired protein for use in medical treatment or in applications using cell culture techniques which require growth factors. The host cell can be a bacterium such as *Bacillus* or *E. coli*, a fungus such as yeast, a plant cell such as tobacco, potato or *arabidopsis* or an animal cell and in particular a vertebrate cell line such as MoCOS or CHO cell lines or an insect cell line.

Yet another subject matter of the present invention is the provision of pharmaceutical compositions which contain a pharmaceutically active amount of a TGF- β -like protein according to the invention as the active substance. If desired, such a composition comprises a pharmaceutically acceptable carrier substance, auxiliary substance, diluent or filler. Such a pharmaceutical composition can be used in wound-healing and tissue regeneration as well as in the healing of damage to bones, cartilage, connective tissue, skin, mucous membranes, epithelium or teeth and in dental implants either alone or in combination with other active substances e.g. other proteins of the TGF- β family or

growth factors such as EGF (epidermal growth factor) or PDGF (platelet derived growth factor). Moreover such a pharmaceutical composition can be used in the prevention of diseases such as the prevention of osteoporosis and arthrosis.

Another possible clinical application of the TGF- β protein according to the invention is to use it as a suppressor of an immunoreaction to prevent rejection of organ transplants or its application in connection with angiogenesis. The pharmaceutical composition according to the invention can also be used prophylactically or in cosmetic surgery. In addition administration of the composition is not limited to humans but can also encompass animals and in particular pets.

Finally the present invention also concerns an antibody that binds specifically to the proteins according to the invention or such an antibody fragment (e.g. Fab or Fab'). Processes for the production of such a specific antibody or antibody fragment are part of the general technical knowledge of an average person skilled in the art. Such an antibody is preferably a monoclonal antibody. Such antibodies or antibody fragments may also be suitable for diagnostic methods.

It is intended to elucidate the invention further by the following example.

Example 1

Isolation of MP-52

1.1 Total RNA was isolated from human embryonic tissue (8 to 9 weeks old) according to the method of Chirgwin et al., Biochemistry 18 (1979), 5294-5299. Poly(A+) RNA was separated from the total RNA by oligo (dT) chromatography according to the manufacturer's instructions (Stratagene Poly (A) Quick columns).

1.2 For the reverse transcription reaction 1 to 2.5 μ g poly (A+) RNA was heated for 5 minutes to 65°C and quickly cooled on ice. The reaction mixture contained 27 U RNA-Guard (Pharmacia), 2.5 μ g oligo (dT)12-18 (Pharmacia), 5 x buffer (250 mmol/l Tris/HCl, pH 8.5, 50 mmol/l MgCl₂, 50 mmol/l DTT, 5 mmol/l of each dNTP, 600 mmol/l KCl) and 20 U AMV reverse transcriptase (Boehringer Mannheim) per μ g poly (A+) RNA. The reaction mixture (25 μ l) was incubated for 2 hours at 42°C.

1.3 The deoxynucleotide primers OD and OID shown in Fig. 2 were prepared on an automatic DNA synthesizer (Biosearch). The purification was carried out by denaturing polyacrylamide gel electrophoresis and isolating the main bands from the gel by means of isotachopheresis. The oligonucleotides were designed by comparing the nucleic acid sequences of known members of the TGF- β family and selecting regions with the highest conservation. A comparison of this region is shown in Fig. 2. In order to facilitate cloning both nucleotides contained EcoRI restriction sites and

OD additionally contained a NcoI restriction site at its 5' terminus.

- 1.4 cDNA corresponding to 20 ng poly (A+) RNA was used as the starting material in the PCR reaction. The reaction was carried out in a volume of 50 μ l and contained 1 x PCR buffer (16.6 mmol/l $(\text{NH}_4)_2\text{SO}_4$, 67 mmol/l Tris/HCl pH 8.8, 2 mmol/l MgCl_2 , 6.7 μ mol/l EDTA, 10 mmol/l β -mercapto-ethanol, 170 μ g/ml bovine serum albumin (Gibco), 200 μ mol/l of each dNTP (Pharmacia), 30 pmol of each oligonucleotide (OD and OID) and 1.5 U Taq polymerase (AmpliTag, Perkin Elmer Cetus). The reaction mixture was overlayed with paraffin and 40 PCR cycles were carried out. The products of the PCR reaction were purified by phenol/chloroform extraction and concentrated by ethanol precipitation.
- 1.5 The PCR reaction product was cleaved with the restriction enzymes SphI (Pharmacia) and AlwNI (Biolabs) according to the manufacturer's instructions.
- 1.6 The products of the restriction cleavage were fractionated by Agarose gel electrophoresis. After staining with ethidium bromide, uncleaved amplification products were cut out of the gel and isolated by phenol extraction. The DNA obtained was subsequently purified twice by phenol/chloroform extraction.

- 1.7 After an ethanol precipitation, a quarter or a fifth of the isolated DNA was reamplified using the same conditions as for the primary amplification except that the number of cycles was reduced to 13. The reamplification products were purified, cleaved with the same enzymes as above and the uncleaved products were isolated from Agarose gels as elucidated above for the amplification products. The reamplification step was repeated twice.
- 1.8 After the last isolation from the gel, the amplification products were cleaved by 4 units EcoRI (Pharmacia) under the conditions recommended by the manufacturer. A fourth of the restriction mixture was ligated into the vector pBluescriptII SK+ (Stratagene) cleaved with EcoRI. 24 clones were analysed further by means of sequencing after ligation. The sample cleaved with AlwNI and SphI resulted in a new sequence that was denoted MP-52. The other clones mainly contained BMP6 sequences and one contained a BMP7 sequence.

The clone was completed up to the 3' end of the cDNA according to the method described in detail by Frohmann (Amplifications, published by Perkin-Elmer Corp., Issue 5 (1990), pp 11-15). The same embryonic mRNA that had been used to isolate the first fragment of MP-52 was reversally transcribed as described above. The amplification was carried out using the adapter primer (AGAATTCGCATGCCATGGTCGACG) and an internal primer (CTTGAGTACGAGGCTTCCACTG) of the MP-52 sequence. The amplification products were reamplified using an overlapping adapter primer (ATTCGCATGCCATGGTCGACGAAG) and an overlapping internal primer (GGAGCCCACGAATCATGCAGTCA) of the MP-52 sequence. After

restriction cleavage with NcoI the reamplification products were cloned and sequenced into a vector that was cleaved in the same way (pUC 19 (Pharmacia No. 27-4951-01) having a modified multiple cloning site which contains a single NcoI restriction site) and sequenced. The clones were characterized by their sequence overlapping at the 3' end of the known MP-52 sequence. One of these was used as a probe to screen a human genomic gene bank (Stratagene No. 946203) according to a method described in detail by Ausubel et al. (Current Protocols in Molecular Biology, published by Greene Publishing Associates and Wiley-Interscience (1989)). One phage (λ 2.7.4) was isolated from 8×10^5 λ phages which contained an insertion of about 20 kb and which is deposited at the DSM under the depository number 7387. This clone contains further sequence informations at the 5' end in addition to the sequence isolated from mRNA by the described amplification methods.

For the sequence analysis a HindIII fragment of about 7.5 kb was subcloned into a vector cleaved in the same manner (Bluescript SK, Stratagene No. 212206). This plasmid denoted SKL 52 (H3) MP12 was also deposited at the DSM under the depository number 7353. The sequence information shown in SEQ ID NO. 1 was derived from the phage λ 2.7.4.. The ATG at position 640 is the first ATG within the reading frame (a stop codon occurs at position 403). Based on the sequence data it may be assumed that this is the start codon for the translation.

The genomic DNA contains an intron of about 2 kb between base pairs 1270 and 1271 of SEQ ID NO: 1. The sequence of the intron is not shown. The correctness of the splicing site was confirmed by sequencing an

amplification product which was derived from cDNA containing this region. These sequence informations were obtained using a slightly modified method which is described in detail by Frohmann (Amplifications, published by Perkin-Elmer Corporation, Issue 5 (1990), pp 11-15). The same embryonic RNA that was also used to isolate the 3' end of MP-52 was reverse transcribed using an internal primer orientated in the 5' direction of the MP-52 sequence (ACAGCAGGTGGGTGGTGTGGACT). A polyA tail was attached to the 5' end of the first cDNA strand using terminal transferase. A two-step amplification was carried out, firstly by using a primer composed of oligo dT and an adapter sequence (AGAATTCGCATGCCATGGTTCGACGAAGC(T16)) and secondly an adapter primer (AGAATTCGCATGCCATGGTTCGACG) and an internal primer (CCAGCAGCCCCATCCTTCTCC) from the MP-52 sequence. The amplification products were reamplified using the same adapter primer and an overlapping internal primer (TCCAGGGCACTAATGTCAAACACG) from the MP-52 sequence. Subsequently the reamplification products were reamplified using an overlapping adapter primer (ATTCGCATGCCATGGTTCGACGAAG) and an overlapping internal primer (ACTAATGTCAAACACGTACCTCTG) from the MP-52 sequence. The final reamplification products were cloned with blunt ends into a vector (Bluescript SK, Stratagene No. 212206) which had been cleaved with EcoRV. The clones were characterized by their sequence overlapping with the DNA of λ 2.7.4..

In addition a cDNA bank - produced from RNA of human fibroblasts and cloned into λ gt10 - was screened. In this process 2×10^6 phages were tested using a ca. 1 kb fragment of genomic MP-52 DNA (2nd exon up to the HindIII restriction site in the 3' untranslated region) as a radioactive probe. 17 mixed plaques were picked out

and were checked by PCR using primers from the 5' and 3' region of the MP-52 sequence. Subsequently 8 phage plaques were selected and isolated. cDNA was isolated from the phage by an EcoRI partial cleavage and cloned into the Bluescript vector that was also cleaved with EcoRI.

Sequencing of one of the resulting plasmids SK52L15.1MP25 showed that the longest phage (15.1) starts at nucleotide No. 321 of SEQ ID NO. 1. In addition the splicing position (nucleotide 1270) was confirmed by the sequencing.

The plasmid SKL 52 (H3) MP12 was deposited on 10th December 1992 at the DSM ("Deutsche Sammlung von Mikroorganismen und Zellkulturen, Mascheroder Weg 1b, 38124 Braunschweig) under the depository number 7353.

The phage λ 2.7.4 was deposited on 13th January 1993 at the DSM under the depository number 7387.

The plasmid SK52L15.1MP25 was deposited on 16th July 1993 at the DSM under the depository number 8421.

Example 2

Expression of MP52

Various systems were checked for the expression of MP52. The use of Vaccinia viruses as an expression system is described in detail and capable of being reproduced by a person skilled in the art in Current Protocols in Molecular Biology (Ausubel et al., Greene Publishing Associates and Wiley-Interscience, Wiley & Sons),

abbreviated CP in the following, in chapter 16 unit 16.15-16.18. The system is based on the fact that foreign DNA can be integrated by homologous recombination into the genome of the Vaccinia virus using certain vectors. For this purpose the vector used contains the TK (thymidine kinase) gene from the Vaccinia genome. In order to enable selection for recombinant viruses the vector in addition contains the E. coli xanthine-guanine phosphoribosyl transferase gene (gpt) (Falkner et al., J. Virol. 62 (1988), 1849-1854). The cDNA with the entire region coding for MP52 was cloned into this vector. The cDNA comes from plasmid SK52L15.1MP25 (DSM, depository number 8421) which was, however, firstly deleted and subcloned in order to remove a large portion of the 5' untranslated region. For this the plasmid SK52L15.1MP25 was linearized with SalI and the 5' end was deleted stepwise with the ExoIII/mung bean kit (Stratagene #200330) according to the manufacturer's instructions. After restriction with BamHI, the MP52 cDNAs that had been deleted to different extents were separated from the residual vector and isolated by an agarose gel and subcloned (pSK52s) according to standard methods (Sambrook et al., Molecular Cloning, second edition, Cold Spring Harbor Laboratory Press 1989) in a pBluescriptII SK vector (Stratagene #212206) restricted with EcoRV and BamHI. All restrictions were carried out according to the manufacturer's instructions. Sequencing with Sequenase (USB/Amersham #70770) yielded inter alia a clone which starts with nucleotide 576 in SEQ ID NO. 1 (64 base pairs distant from the start codon). The cDNA insert was isolated from this by restriction with SalI and SacI and cloned into the likewise cleaved vector for recombination in the vaccinia virus system. The resulting plasmid (pBP1MP52s) was deposited on 24th May

1994 at the DSM (deposit number 9217) and used for the production of recombinant Vaccinia viruses. For this up to 80 % confluent 143B cells (HuTk, ATCC CRL 8303) in 35 mm culture dishes were infected with Vaccinia wild-type virus in 2 ml PBS for 30 minutes at room temperature while shaking occasionally (1 virus per 10 cells). After aspirating the supernatant and adding 2 ml culture medium (MEM, Gibco BRL #041-01095), it was incubated for 2 hours at 37°C. The medium was subsequently removed and transformation of these cells was achieved with 100 ng pBP1MP52s, 2 µg carrier DNA (calf thymus, Boehringer Mannheim #104175) and 10 µl Lipofectin (Gibco BRL #18292-011) in 1 ml MEM for 15 hours at 37°C. After addition of 1 ml MEM containing 20 % FCS (Gibco BRL #011-06290), they were incubated for a further 24 hours at 37°C and subsequently the lysed cells were frozen.

The gpt selection for xanthine-guanine phosphoribosyl transferase and isolation and amplification of individual recombinant viruses was essentially carried out as described in unit 16.17 of CP except that RK13 cells (ATCC CCL 37) were used.

Integration of MP52 cDNA into the virus genome was confirmed by dot blot and Southern blot analysis (CP unit 16.18). A recombinant virus was used for expression analyses in the cell line 143B (HuTk-, ATCC CRL 8303, human). Confluent cells were infected for 45 minutes at 37°C with a number of viruses corresponding to the number of cells and subsequently added to the respective culture medium (MEM, Gibco BRL #041-01095) containing 10 % FCS and penicillin/streptomycin (1:500, Gibco BRL #043-05140H). After 6 hours at 37°C, the medium was removed, the cells were washed twice with e.g. HBSS (Gibco BRL #042-04180M) and production medium (e.g. MEM)

without FCS was added. After 20 to 22 hours of production the cell supernatant was collected. The expression was analysed by means of Western blots according to standard methods (CP unit 10.8). For this the proteins from 100 to 500 μ l cell culture supernatant were precipitated by addition of an equivalent volume of acetone and incubating for at least one hour on ice and centrifuged. After resuspending the pellet in application buffer (7 M urea, 1 % SDS, 7 mM sodium dihydrogen phosphate, 0.01 % bromophenol blue and if desired 1 % β -mercaptoethanol) separation was carried out in 15 % polyacrylamide gels. A prestained protein molecular weight standard (Gibco BRL #26041-020) was used as the marker proteins. Transfer onto a PVDF membrane (Immobilon #IPVH00010) and blocking the membrane were carried out according to standard methods.

In order to detect MP52 on the membrane, polyclonal antibodies against MP52 had been produced in chickens as well as in rabbits. For this the mature part of MP52 with 6 histidines at the N-terminus was expressed in *E. coli* and purified as described for example in Hochuli et al. (BIO/Technology, Vol. 6, 1321-1325 (1988)). Both antibodies enable the specific detection of expression of MP52 wherein dimeric MP52 is less efficiently recognized than monomeric. Chicken antibodies were used for the Western blot in Figure 3 that had been specifically purified by means of PEG precipitation (Thalley et al., BIO/Technology vol. 8 934-938 (1990)) and by means of membrane-bound antigen (mature MP52 containing 6 histidines) (18.17 in Sambrook et al., Molecular cloning, second edition, Cold Spring Harbor Laboratory Press 1989). Anti-chicken IgG with coupled alkaline phosphatase (Sigma A9171) was used as the second antibody. The detection was carried out using the

Tropix Western-Light protein detection kit (Serva #WL10RC) according to the manufacturer's instructions.

The Western blot in Figure 3 shows that MP52-specific bands only occur with the recombinant viruses but not with the wild-type viruses (without integrated foreign-DNA). The expression of MP52 results in a secreted protein having an apparent molecular weight of about 25 kDa in the gel under non-reducing conditions. The protein migrates in the gel at 14 to 15 kDa under reducing conditions. These results show that MP52 is expressed as a dimeric mature protein. The weak bands appearing in the region above 60 kDa that occur in the Western blot are probably residues of the uncleaved precursor proteins. The migration properties also confirms the theoretical molecular weights that can be derived from SEQ ID NO. 2 according to which mature, monomeric MP52 has a size of 13.6 kDa.

It has been proven to be possible to express MP52 and cleave the precursor protein to mature MP52 in various cell lines. C127 (ATCC CRL 1616, mouse), BHK21 (ATCC CCL 10, hamster), MRC-5 (ATCC CCL 171, human) and 3T6-Swiss albino (ATCC CCL 96, mouse) cells were tested.

Expression and cleavage to form mature MP52 was also demonstrated in a further eukaryotic expression system. For this cDNA from MP52 (starting with nucleotide 576) was cloned into the expression plasmid pSG5 (Stratagene #216201). The plasmid pSK52s was restricted with ClaI and XbaI and the protruding ends of the MP52 insert were made blunt by T4 polymerase treatment. Cloning into the vector pSG5, that had been restricted with EcoRI and likewise blunt ended by T4 polymerase treatment, was

carried out according to standard methods. All enzymatic reactions were carried out according to the instructions of the manufacturer. Correct orientation of the MP52 insert was ensured by restriction analysis and sequencing with the T7 primer (Stratagene #300302). The resulting plasmid pSG52s (deposited on 17.05.94 at the DSM with the deposit number DSM 9204) can be cotransformed with a vector that codes for a selectable marker such as e.g. the gene for G418 resistance in order to obtain stable cell lines. For this purpose pSG52s was cotransformed with the plasmid p3616 (deposited on 17.05.94 at the DSM with the deposit number DSM 9203) in L929 cells (ATCC CCL1, mouse) using Lipofectin (Gibco BRL #18292-011) according to the manufacturer's instructions. Selection with G418 was carried out according to methods familiar to a person skilled in the art (CP, unit 9.5) and it resulted in a cell line that produced detectable mature MP52 in the Western blot.

A further expression vector for MP52 was produced using the plasmid pABWN (Niwa et al., Gene 108 (1991), 193-200 and Figure 4) which was provided by Dr. Miyazaki.

For this the HindIII fragment from plasmid pSK52s that starts with nucleotide 576 in SEQ ID NO. 1, was isolated and the protruding ends were made blunt by treatment with Klenow fragment. A Not I restriction cleavage site was introduced at both ends of the fragment by ligation of the adapter.

Adapter: AGCGGCCGCT
TCGCCGGCGA

Vector pABWN was restricted with XhoI, also treated with the Klenow fragment and dephosphorylated with intestinal alkaline phosphatase from the calf (Boehringer

Mannheim). The same phosphorylated adapter was ligated on so that an insertion of the MP52 fragment after restriction with NotI into the generated Not I cleavage site of the vector was now possible. The expression vector that results is subsequently denoted HindIII-MP52/pABWN. All the reactions carried out for the cloning were carried out according to standard methods (e.g. CP unit 3.16).

The structure of the HindIII-MP52/pABWN expression vector was confirmed by sequencing and restriction mapping. HindIII-MP52/pABWN contains the MP52 sequence starting with nucleotide 576 and ending with nucleotide 2278 in SEQ ID NO. 1.

HindIII-MP52/pABWN was transfected in L cells* (mouse fibroblasts) and stable transformants were established therefrom.

For this 4 µg in each case of the plasmids (HindIII-MP52/pABWN or pABWN) were transfected in 5×10^5 L cells in a 6 cm culture dish using 20 µl lipofectAMINE reagent (Gibco BRL #18324-012). For this solution A (4 µg of the respective DNA in 200 µl OPTI-MEM I (Gibco BRL #31985)) was carefully mixed with solution B (20 µl lipofectAMINE reagent in 200 µl OPTI-MEM I) and incubated for 45 minutes at room temperature to form the DNA liposome complex. In the course of this the cells were washed once with 2 ml OPTI-MEM I. For each transfection, 1.6 ml OPTI-MEM I was added to the vessel with the DNA liposome complex. The solution was carefully mixed and the washed cells were overlayed therewith. The cells were incubated with the dilute complex for 5 hours at 37°C in an CO₂ incubator. After the incubation 2 ml DMEM (Gibco BRL, Dulbecco's modified eagle medium)/ 20 % FCS was added.

24 hours after the transfection, the medium was replaced with fresh DMEM/ 10 % FCS. 48 hours after the start of transfection, the cells were transferred into a 10 cm culture dish. 72 hours after the start of the transfection, the G418 selection was started at a concentration of 800 µg/ml. The stable clones appeared after 1 to 2 weeks.

5 ml conditioned DMEM with or without FCS was obtained from confluent transformants which had been grown for 3 days in a 10 cm culture dish. The two different cell culture supernatants (HindIII-MP52/pABWN and pABWN) of transfected cells as well as cell lysates were examined by Western blot. In this process mature MP52 was found in conditioned medium as well as in cell lysates of cells transfected with HindIII-MP52/pABWN. The clones were further cloned and cells producing MP52 were each selected after Western blot analysis. Estimations from Western blot analyses yielded a MP52 production of up to 1 mg/l.

Example 3:

Biological activity of MP52

Several experiments were carried out in vitro and in vivo in order to detect the biological activity of MP52 and to prove the usefulness of this invention for medical applications for the prevention and/or treatment of bone diseases.

1. In vitro assays

1.1

Since an increase in glycosaminoglycan (GAG) synthesis is described in chondrocytes after stimulation with TGF- β (Hiraki et al., Biochimica et Biophysica Acta 969 (1988), 91-99), it was examined whether MP52 also has this influence. The chondrogenic activity of MP52 was tested in primary cultures from foetal rat extremities using the cell culture supernatants (DMEM containing 10 % FCS) of L cell transformants producing MP52 (transfected with HindIII MP52/pABWN).

The four extremities of 16-day-old rat fetuses were used for this. After trypsinization, the cells obtained in F-12 medium (Nutrient mixture Ham's F-12, Gibco BRL #21700) containing 10 % FCS were plated out at 3×10^5 cells on 24-well plates coated with collagen type I and cultured for ca. 2 days until confluence. 56 μ l conditioned medium (CM) of HindIII-MP52/pABWN-L cell transfectants, of pABWN-L cell transfectants or only medium (DMEM containing 10 % FCS) was added to 500 μ l culture medium in each case (F-12 medium containing 10 % FCS). F-12 medium containing 10 % FCS as well as the respective additives was used over a period of 0, 3, 6 and 9 days. The medium containing the respective additives was exchanged every three days. Afterwards the culture was cultured for a further 2 days in F-12 medium without FCS in the presence of the respective additives (conditioned medium or control medium) and then ^{35}S sulfate was added for 6 hours. ^{35}S incorporated into polysaccharides was measured after pronase E digestion

and precipitation as described in Hiraki et al.
(Biochimica et Biophysica Acta 969 (1988), 91-99).

Table 1:

Number of days of incubation	Radioactivity (cpm/well)		
	DMEM (10 % FCS) from control L cells	CM from pABWN-L cell transfectants	CM from HindIII- MP52/pABWN-L cell transfectants
2	3720±114	3865±120	4879±422
5	4188±135	4154±29	8223±275*
8	3546±160	3310±115	9890±1260*
11	3679±218	3633±167	7520±160*

Values relate to \pm S.E.M. for 3 or 4 culture mixtures
*: $p < 0.01$ vs DMEM and CM from pABWN-L cell transfectants
(Scheffe's multiple t-test)

As shown in Table 1, the cell culture supernatants of the transfectants producing MP52 significantly stimulate GAG synthesis in comparison to pure culture medium (DMEM containing 10 % FCS) or to the cell culture supernatant from L cells transfected with pABWN. This shows that MP52 can stimulate differentiation of chondrocytes.

1.2

An effect which has been described for some members of the BMP family is the stimulation of alkaline phosphatase (ALP) activity in osteoblasts. The clonal rat cell line ROB-C26 (C-26) is among the osteoblasts at a relatively early stage of maturation (Yamaguchi et al, Calcif. Tissue Int. 49 (1991), 221-225). The capability of stimulating ALP activity is described for osteoinductive proteins such as e.g. BMP-2 by Yamaguchi et al. (J. Cell Biol. 113 (1991), 681-687).

The influence of MP52 on C26 cells was examined as follows: C26 cells were plated out at 3×10^4 cells per well in a 24-well plate and cultured in α -MEM (Gibco BRL) / 10 % FCS until confluence. 56 μ l of the cell culture supernatant from L cell transfectants producing MP52 (Hind III-MP52/pABWN) or of the cell culture supernatant from pABWN-L cell transfectants or only of the cell culture supernatant (DMEM containing 10 % FCS) from L cells was added to 500 μ l of the C-26 cell culture medium. A change of medium with the respective additives was carried out every three days. The ALP activity in the cell extracts was determined after 0, 3, 6, 9 and 12 days with the aid of standard techniques based on p-nitrophenyl phosphate as the substrate as described for example by Takuwa et al. (Am. J. Physiol. 257 (1989), E797-E803).

Table 2:

ALP activity (nmol/min) per well			
Number of incubation days	DMEM (10 % FCS) from control L cells	CM from pABWN L cell transfectants	CM from HindIII-MP52/pABWN-L cell transfectants
0	41.8±2.8	41.8±2.8	41.8±2.8
3	136.3±3.7	125.8±2.3	181.3±14.2*
6	129.0±7.8	119.3±6.4	258.0±8.3*
9	118.4±3.7	110.1±2.8	258.4±10.6*
12	121.2±3.2	125.3±6.0	237.8±11.0*

Values relate to \pm S.D. for 4 culture mixtures.

*:p<0.01 vs DMEM and CM from pABWN-L cell transfectants (Scheffe's multiple t-test)

As shown in Table 2, addition of MP52 leads to a significant increase in ALP activity compared to pure DMEM/10 % FCS medium and medium from pABWN-infected L cells. This result shows that MP52 cannot only cause chondrocytes to differentiate but can also lead to the differentiation and maturation of osteoblasts.

A further osteoblast cell line (MC3T3-E1, mouse) that shows an increase in the ALP activity by treatment with BMP-2 as described by Takuwa et al. (Biochem. Biophys. Res. Com. 174 (1991), 96-101), does not result in any

change in the ALP activity after incubation with conditioned medium from L cell transfectants producing MP52 (HindIII-MP52/pABWN) or medium after MP52 production by infection with recombinant Vaccinia viruses. This indicates that MP52 has a cell specificity that partially deviates from that of BMP-2. Different functions due to different target sites for the individual TGF- β family members may be of great medical relevance.

2. In vivo experiments

2.1

The most definitive possibility of examining bone development is based on ectopic bone formation in vivo. This can for example be induced by implantation of demineralized bone matrix (Urist, Science 150 (1965), 893-899). The same process can be induced by combination of inactive matrix with bone-inducing proteins as described for example by Sampath et al. (PNAS* Proc.Natl.Acad.sci. USA 78 (1981), 7599-7603). This process of bone formation is similar to embryonic enchondral bone formation and adult bone healing. This method therefore enables proteins to be examined for their bone-inductive capability in vivo.

MP52 protein which had been obtained by expression in the Vaccinia system (see example 2) was partially purified and implanted for such an experiment.

For this 143B cells (HuTk, ATCC CRL 8303) were cultured in culture dishes and roller flasks until confluence and infected with recombinant viruses as described in

example 2 for the expression analyses, they were washed and MP52 was allowed to accumulate for about 20 hours in MEM (Gibco BRL, ca 1 ml per 10^6 cells). The same preparation was infected with wild-type viruses as a control. Cell culture supernatant (conditioned medium) from each preparation was collected and centrifuged ($40000 \times g$ for 30 minutes at 4°C). In order to remove the viruses, the supernatants were filtered over inorganic filters ($0.1 \mu\text{m}$ pore size, Whatman, Anotop 25). In the course of the characterization of MP52 it was shown that this protein binds to heparin-Sepharose. This property was utilized for partial purification. For this the filtered and centrifuged, conditioned medium was brought to a final concentration of 50 mM Tris pH 7.0, 100 mM NaCl and 6 M urea and it was loaded onto a heparin column (HiTrapTM, Pharmacia #17-0407-01) that uses equilibrated in buffer A (50 mM Tris pH 7.0, 100 mM NaCl and 6 M urea). The loaded column was washed with buffer A and eluted with a linear gradient to 100 % buffer B (50 mM Tris pH 7.0, 600 mM NaCl and 6 M urea) at a flow rate of 0.5 ml/min within 50 min (2.5 ml per fraction). The use of urea is not absolutely necessary. MP52 elutes reproducibly mainly in 2 fractions at about 250 to 400 mM NaCl as could be examined by Western blot analysis (see example 2). Aliquots of these fractions were also examined in 15 % polyacrylamide gels stained with silver according to the instructions of the manufacturer (Silver Stain-II, Daiichi #SE140000) and the fractions were pooled. The comparable fractions were also pooled after analysis in gels stained with silver after purification from conditioned medium after infection with wild-type viruses.

Further examinations on MP52 showed that MP52 also binds to hydroxyapatite. Thus it is in principle possible to

achieve an additional purification by a hydroxyapatite column or to replace a heparin column by a hydroxyapatite column (e.g. BIO-RAD, Econo-pac HTP). Other methods known to a person skilled in the art are also conceivable for further purifications such as e.g. gel sieve columns, ion exchanger columns, affinity columns, metal chelate columns or columns based on hydrophobic interactions.

The MP52 protein prepurified by heparin-Sepharose chromatography or the corresponding proteins that are still contaminated which are also present in the cell culture supernatants infected with the wild-type, were further purified by means of reversed phase HPLC. For this a C8 column (Aquapore RP300, Applied Biosystems, particle size: 7 μm , pore size: 300 Å) was equilibrated with 10 % buffer B (buffer A: 0.1 % trifluoroacetic acid; buffer B: 90 % acetonitrile, 0.1 % trifluoroacetic acid). After loading the column with the pooled fractions containing MP52 from the heparin column it was extensively washed with 10 % buffer B. The bound protein was eluted with the following gradient: 10 to 50 % buffer B for 20 minutes and 50 to 100 % buffer B for 50 minutes. Fractions of 500 μl were collected and analysed by Western blot as well as with gels stained with silver. The MP52 protein elutes under the selected conditions in the range of about 55 to 65 % acetonitrile. The fractions containing MP52 were pooled. The same procedure was carried out with the corresponding fractions from the control purification of cell culture supernatant from cells infected with wild-type viruses.

Partially purified MP52 protein at a concentration estimated to be 50 ng/ml according to Western blot

analysis also showed a significant increase in the ALP activity in ROB-C26 cells after three days of incubation.

Partially purified MP52 protein or control protein from the corresponding partially purified cell culture supernatants after infection with wild-type viruses were reconstituted with matrix and implanted in rats in order to prove its capability for cartilage and bone formation.

In principle various matrix materials known to a person skilled in the art should be usable i.e. natural (also modified) and synthetically prepared matrices, however, biocompatible porous materials that can be biologically degraded are preferred. In these experiments bone matrix from rats was used that had been prepared essentially in a similar way to that described by Sampath et al. (PNAS 80 (1983), 6591-6595). The rat bones (femur and tibia) were demineralized in 0.6 M HCl for 24 hours and subsequently bone marrow that was still present was removed. After washing with water and defatting for three hours in a chloroform/methanol (1/1) mixture, the bones were air-dried, powderized in a mill in a deep-frozen state and particle sizes between 400 and 1000 μm were sieved out. Subsequently the matrix was extracted for 7 days at room temperature in 4 M guanidinium HCl in the presence of protease inhibitors. After washing extensively with water, the matrix was lyophilized and stored at 4°C. Matrices treated in this way do not on their own show bone-inducing activity.

Protein can be combined with the extracted bone matrix by various methods known to a person skilled in the art.

MP52 protein or control protein that had been purified by means of heparin-Sepharose as well as by reversed phase HPLC, was combined after elution in the acetonitrile/trifluoro-acetic acid solution with 25 mg matrix in each case per implant, mixed well, deep-frozen and lyophilized.

For the implantation of matrix-bound MP52, two ca. 3 months-old rats (Whistar) were used which had been anaesthetised by intramuscular injection of an anaesthetic (0.2 ml Rompun (Bayer) mixed with 0.5 ml Ketanest 50 (Parke Davis)) using 0.14 ml per 100 g body weight. Bilateral pockets were prepared in the abdominal muscles for the implants (beneath the thorax, starting ca. 0.5 cm below the lowest costal arch). The matrix-bound MP52 (ca. 2 to 4 μ g as estimated by Western blot) as well as the corresponding matrix-bound control proteins were moistened using 0.9 % saline solution (Delta Pharma) and introduced into the muscular pockets. The muscular pockets as well as the necessary skin incisions were subsequently sutured. The rats were immunosuppressed with Cyclosporin A (Sandimmun).

After 18 or after 26 days the implants were removed from the rats and fixed for histological examinations. Since after 26 days the implant with MP52 allowed the assumption that macroscopically bone formation had already occurred, this was embedded in methylmethacrylate in order to prepare thin sections, the other implants were embedded in paraffin. Mineralized cartilage and bone tissues are accentuated in black by means of the von Kossa staining technique (Romeis, B.; "Mikroskopische Technik", Ed: Böck, P.; Urban and Schwarzenberg; Munich, Baltimore, Vienna (1989)). In the trichromium staining according to Masson-Goldner

(Romeis, B.; "Mikroskopische Technik", Ed: Böck, P.; Urban and Schwarzenberg; Munich, Baltimore, Vienna (1989)), mineralized bone tissue and collagen are stained bright green, osteoid is stained red and cytoplasm reddish-brown. Both staining techniques were used on implants from both rats. In both experimental animals considerable formation of cartilage and bone was detected in the implants containing MP52 using both staining techniques. The corresponding implants with control protein showed no formation whatsoever of cartilage or bone. The number of cartilage precursors with chondrocytes and cartilage areas with initial formation of extracellular matrix and its mineralization in concentric circles is higher in the MP52 implant after 18 days than in the one after 26 days. Mature bone tissue with vectorial osteoid formation as well as individual osteocytes in the bone is, however, also detectable in the implant after 18 days. In addition closed ossicles can be observed with the onset of bone marrow formation. In the implant after 26 days areas of cartilage with initial matrix formation and calcification are also detectable, the portion of mineralized bone tissue stained green and having osteocytes and osteoid edges has, however, significantly increased. In this implant bone marrow formation together with the occurrence of isolated fat cells can also be detected. For illustration Figure 5 shows the staining test of the bone material (according to von Kossa stain.) from the entire implant after 26 days. A small section of the same implant is shown in Figure 6 after staining according to Masson-Goldner. It shows active bone with an edging of cuboidal osteoblasts and osteoid in which individual embedded osteoblasts can be recognized. Furthermore individual osteocytes can also be seen in the mineralized bone tissue (stained green in

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the original preparation). The formation of bone marrow is also detectable.

The experiment shows that recombinantly produced MP52 alone, in combination with a matrix is capable of inducing enchondral bone formation.

2.2

In order to confirm the results, a further ectopic test for bone formation using MP52 L cell transformants was carried out. L cells (1×10^6 cells) producing MP52 (transfected with HindIII-MP52/pABWN) and non-producing (pABWN-transfected) L cells were injected into the bilateral thigh muscles of three male naked mice in each case. All animals were killed after three weeks, the thigh muscles were excised and these were examined with low energy X-ray radiation as well as histopathologically.

Analysis by X-ray radiation shows dense material at the injection sites in the muscle tissue of all L cells producing MP52 as listed in Table 3. Simple cartilage formation and calcified cartilage formation could be determined in the muscles using histological examinations. These results also confirm that MP52 can induce enchondral bone formation.

Table 3:

	Cells producing MP52 (HindIII- MP52/pABWN)	Control cells (pABWN)
dense material by X-ray analysis	3/3	0/3
chondrocytes by histology	3/3	0/3
calcified cartilage formation by histology	3/3	0/3

The experiments that were carried out confirm that MP52 protein stimulates the formation of cartilage from undifferentiated mesenchymal cells as well as the differentiation and maturation of osteoblasts. This leads to enchondral bone formation which is similar to the induction cascade in embryonic bone formation and bone healing of fractures.

The conditions stated in the experiments are to be looked upon as an illustration of the MP52 activity but not as limitation. The invention can also be examined and characterized in another form.

CLAIMS

1. DNA molecule that codes for a protein of the TGF- β family and which comprises
 - (a) the part coding for the mature protein and if desired further functional parts of the nucleotide sequence shown in SEQ ID NO. 1,
 - (b) a nucleotide sequence corresponding to the sequence from (a) within the scope of the degeneracy of the genetic code,
 - (c) a nucleotide sequence corresponding to an allelic derivative of one of the sequences from (a) and (b), or
 - (d) a nucleotide sequence hybridizing with one of the sequences from (a), (b) or (c) provided that a DNA molecule according to (d) completely contains at least the part coding for a mature protein of the TGF- β family.
2. Vector,
w h e r e i n
it contains at least one copy of a DNA molecule as claimed in claim 1.
3. Host cell,
w h e r e i n
it is transformed by a DNA as claimed in claim 1 or by a vector as claimed in claim 2.

4. Host cell as claimed in claim 3,
w h e r e i n
it is a bacterium, a fungus, a plant or an animal
cell.
5. Protein of the TGF- β family which is coded by a DNA
sequence as claimed in claim 1.
5. Protein as claimed in claim 5,
w h e r e i n
it has the amino acid sequence shown in SEQ ID
NO. 2 or, if desired, functional parts thereof.
7. Process for the production of a protein of the
TGF- β family,
w h e r e i n
a host cell as claimed in claim 3 or 4 is cultured
and the TGF- β protein is isolated from the cell
or/and from the culture supernatant.
8. Pharmaceutical composition,
w h e r e i n
it contains at least one protein as claimed in
claim 5 or 6 as the active substance if desired,
together with the usual pharmaceutical carrier
substances, auxiliary substances, diluents or
fillers.
9. Pharmaceutical composition as claimed in claim 8
for the treatment or prevention of damage to bone,
cartilage, connective tissues, skin, mucous
membranes, epithelium or teeth, for application in
dental implants and for application in wound-
healing and tissue regeneration processes.

10. Antibody or antibody fragments,
w h e r e i n
they bind to a protein as claimed in claim 5 or 6.

ABSTRACT

The invention concerns a protein of the TGF- β family,
the DNA coding therefor and a pharmaceutical composition
containing the protein.

-9. Juni 1994

Figure 1

	10	20	30	40	50	
MP 52	CSRKALEVNF	KMGWDDWII	APLEYEAFBC	BGLCEPPLRS	BLEPTNEAVI	
BMP 2	CRREPLYVDF	SDVGNDWIV	APPGYDAFYC	BGECPPLAD	HINSTINEAIV	
BMP 4	CRRESLYVDF	SDVGNDWIV	APPGYDAFYC	BGECPPLAD	HINSTINEAIV	
BMP 5	CKKBELVVSF	RDLGWQDWII	AFEGYDAFYC	DGECSFPLNA	EMNATINEAIV	
BMP 6	CKKBELVVSF	QDLGWQDWII	AFKGYDAFYC	DGECSFPLNA	EMNATINEAIV	
BMP 7	CKKBELVVSF	RDLGWQDWII	AFEGYDAFYC	BGECPPLRS	YMNATINEAIV	
	* +	* * *	* * * * * +	**	* * +	* + * * * + + * * *
	60	70	80	90	100	
MP 52	QILVNSDPE	SIPPTCCVPT	RLSPISILFI	DSNNVVYKQ	YEDMVVESCOG	CR
BMP 2	QILVNSVNS-	KIPKACCVPT	ELSAISMXYL	DEYKVVVLN	YQDMVVESCOG	CR
BMP 4	QILVNSVNS-	SIPKACCVPT	ELSAISMXYL	DEYKVVVLN	YQDMVVESCOG	CR
BMP 5	QILVHMFDP	YVPKPCCAPT	KINRISVLXF	DSSNVILNK	YRNMVVRSOG	CH
BMP 6	QILVHMFDP	YVPKPCCAPT	KINRISVLXF	DSSNVILNK	YRNMVVRSOG	CH
BMP 7	QILVHMFDP	YVPKPCCAPT	QINRISVLXF	DSSNVILNK	YRNMVVRSOG	CH
	*** + + +	+ + + *	** + + +	* + * * *	* + + + *	* + * * * + + + + * +

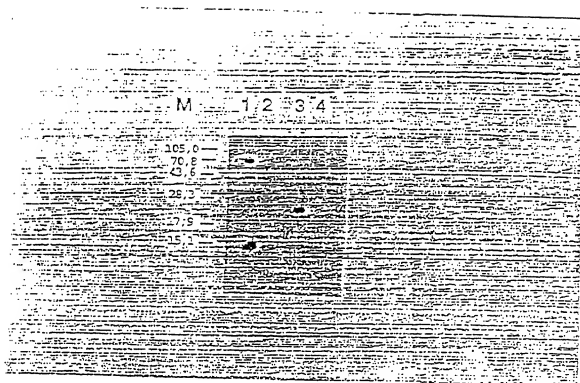
Figure 2a

	Eco RI Nco I
OD	ATGAATTCOCATGCACTGGGCTGG-ATGATTCGAT
BMP 2	ACCTGGGCTGGCAATGACTGGAT
BMP 3	ATATTCGGCTGGAGTGAATGGAT
BMP 4	ATGTGGGCTGGCAATGACTGGAT
BMP 7	ACCTGGGCTGGCAGGACTGGAT
TGF-β1	AGGACCTGGGCTGGCAATGGAT
TGF-β2	GGGATCTAGGCTGGCAATGGAT
TGF-β3	AGGATCTGGGCTGGCAATGGAT
Inhibin α	AGCTGGGCTGGCAAGCTGGAT
Inhibin β _A	ACATCGGCTGGCAATGACTGGAT
Inhibin β _B	TCATCGGCTGGCAAGACTGGAT

Figure 2b

	Eco RI
OID	ATGAATTCGAGCTGGGTGGGTCACAGCA
BMP 2	GAGTCTCTGGGTCACAGCA
BMP 3	CATCTTTTCTGGTACACAGCA
BMP 4	CAGTTCAGTGGGTCACAGCA
BMP 7	GAGCTGGGTGGGTCACAGCA
TGF-β1	CAGGGCTGGGTCAGGAGCA
TGF-β2	TAAATCTGGGTCAGGAGCA
TGF-β3	CAGGTCTGGGTCAGGAGCA
Inhibin α	CCCTGGGAGAGTACACAGCA
Inhibin β _A	CAGCTTGGTGGGTCACAGCA
Inhibin β _B	CAGCTTGGTGGGTCAGCA

Figure 3



M: prestained protein molecular weight marker with the stated apparent molecular weights listed (Gibco BRL #26041-020)

- 1: Cell culture supernatant (100 μ l) after infection with recombinant viruses (with inserted MP52 cDNA) under reducing (1 % β -mercaptoethanol) conditions
- 2: Cell culture supernatant (100 μ l) after infection with wild-type viruses (without inserted foreign DNA) under reducing (1 % β -mercaptoethanol) conditions
- 3: Cell culture supernatant (500 μ l) after infection with recombinant viruses (with inserted MP52 cDNA) under non-reducing conditions
- 4: Cell culture supernatant (500 μ l) after infection with wild-type viruses (without inserted foreign DNA) under non-reducing conditions

Figure 4

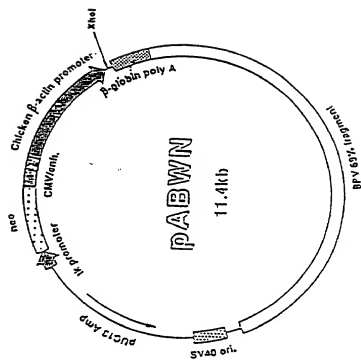




Figure 5

Section through the entire implant (26 days after implantation) stained according to Kossa. Mineralized tissue in black is clearly distinguishable from the surrounding muscle tissue.

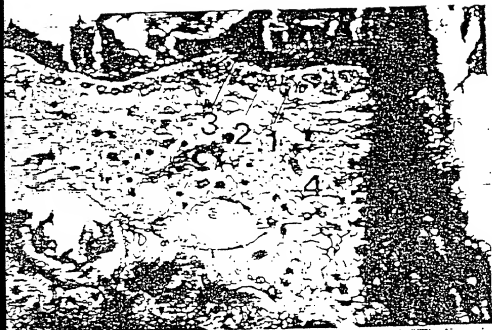


Figure 6

Cross-section of the implant (26 days after implantation) stained according to Masson-Goldner.

1: border consisting of osteoblasts (pink in the original)

2: osteoid (red in the original)

3: mineralized bone tissue (green in the original) with osteocytes
(pink in the original)

4: bone marrow (slight pink to orange in the original)

Fig.1

Fig.1a
Fig.1b

Fig.1a

	10	20	30	40	50
SEQ ID NO 13 MP 52 CSRKALHVNF	KDMGWDDWII	APLEYEAFHC	EGLCEFP LRS	HLEPTNHA IV	
SEQ ID NO 14 BMP 2 CKRHPLYVDF	SDVGWNDWIV	APPGYHAFYC	HGECFPPLAD	HLNSTNHA IV	
SEQ ID NO 15 BMP 4 CRRHSLYVDF	SDVGWNDWIV	APPGYQAFYC	HGDCFPPLAD	HLNSTNHA IV	
SEQ ID NO 16 BMP 5 CKKHLYVVSF	RDLGWQDWII	APEGYAAFYC	DGECSFPLNA	HMNATNHA IV	
SEQ ID NO 17 BMP 6 CRKHLYVVSF	QDLGWQDWII	APKGAAANYC	DGECSFPLNA	HMNATNHA IV	
SEQ ID NO 18 BMP 7 CKKHLYVVSF	RDLGWQDWII	APEGYAAFYC	EGECAPPLNS	YMNATNHA IV	
	* + * * *	** * * *	++ * * * *	++ * * *	***

Fig.1b

	60	70	80	90	100
SEQ ID NO 13 MP 52 QTLNMSDPE	STPPTCCVPT	RLSPISILFI	DSANNVVYKQ	YEDMVVESCQ	CR
SEQ ID NO 14 BMP 2 QTLVNSVNS-	KIPKACCVPT	ELSAISMLYL	DENEKVVILKN	YQDMVVEGCG	CR
SEQ ID NO 15 BMP 4 QTLVNSVNS-	SIPKACCVPT	ELSAISMLYL	DEYDKVVILKN	YQDMVVEGCG	CR
SEQ ID NO 16 BMP 5 QTLVHLMFPD	HVPKPCCAPT	KLNAISVLYF	DDSSNVI LKK	YRNMVVRSCG	CH
SEQ ID NO 17 BMP 6 QTLVHLMNPE	YVPKPCCAPT	KLNAISVLYF	DDNSNVI LKK	YRNMVVRACG	CH
SEQ ID NO 18 BMP 7 QTLVHFINPE	TVPKPCCAPT	QLNAISVLYF	DDSSNVI LKK	YRNMVVRACG	CH
*** +++ ++	+ + ***+	++ ** *	* *** *	* +*****	**

Fig.2a

Eco RI Nco I

SEQ ID NO 19 OD	ATGAATTC	CCCATGGACCTGGGCTGGMAKGAMTGGAT
SEQ ID NO 20 BMP 2		ACGTGGGGTGGAAATGACTGGAT
SEQ ID NO 21 BMP 3		ATATTGGCTGGAGTGAATGGAT
SEQ ID NO 22 BMP 4		ATGTGGGCTGGAATGACTGGAT
SEQ ID NO 23 BMP 7		ACCTGGGCTGGCAGGACTGGAT
SEQ ID NO 24 TGF- β 1		AGGACCTCGGCTGGAAGTGGAT
SEQ ID NO 25 TGF- β 2		GGGATCTAGGGTGGAAATGGAT
SEQ ID NO 26 TGF- β 3		AGGATCTGGGCTGGAAGTGGGT
SEQ ID NO 27 INHIBIN α		AGCTGGGCTGGGAACGGTGGAT
SEQ ID NO 28 INHIBIN β_A		ACATCGGCTGGAATGACTGGAT
SEQ ID NO 29 INHIBIN β_B		TCATCGGCTGGAACGACTGGAT

Fig.2b

Eco RI

SEQ ID NO 30 OID	ATGAATTC	GAGCTGCGTSGGSRACACAGCA
SEQ ID NO 31 BMP 2		GAGTTCTGTGGGACACACAGCA
SEQ ID NO 32 BMP 3		CATCTTTTCTGGTACACACAGCA
SEQ ID NO 33 BMP 4		CAGTTCAGTGGGCACACAACA
SEQ ID NO 34 BMP 7		GAGCTGCGTGGGCACACAGCA
SEQ ID NO 35 TGF- β 1		CAGCGCCTGCGGCACGCAGCA
SEQ ID NO 36 TGF- β 2		TAAATCTTGGGACACGCAGCA
SEQ ID NO 37 TGF- β 3		CAGGTCTTGGGACACGCAGCA
SEQ ID NO 38 INHIBIN α		CCCTGGGAGAGCAGCACAGCA
SEQ ID NO 39 INHIBIN β_A		CAGCTTGGTGGGCACACAGCA
SEQ ID NO 40 INHIBIN β_B		CAGCTTGGTGGGAATGCAGCA

Fig.3

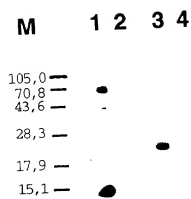


Fig.4

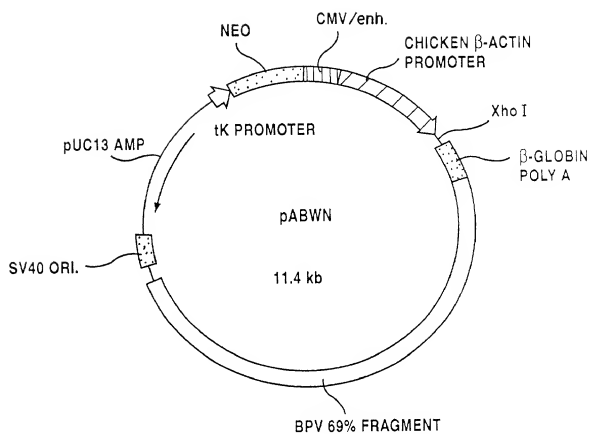
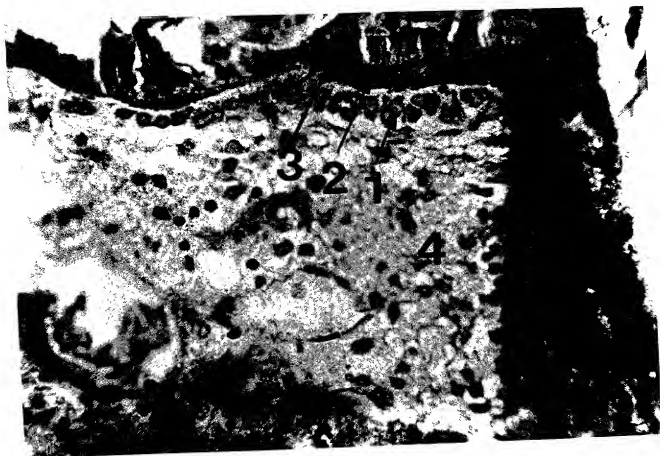


Fig.5



09264501002100

Fig.6



001200 051800

Declaration For U.S. Patent Application

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled
(Insert Title) NEW GROWTH/DIFFERENTIATION FACTOR OF THE TGP-B FAMILY
the specification of which

- (Check one of blocks 1, 2 or 3. See note A on back of this page)
1. ☐ is attached hereto.
 2. ☐ was filed on _____ as International PCT Application Serial No. _____ and was amended on _____ (if applicable)
 3. ☒ was filed on August 10, 1994 as U.S. Application Serial No. 08/288,508 and was amended on _____ (if applicable)

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claim(s), as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, §1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application for which priority is claimed:

(List prior foreign applications. See note B on back of this page)	<u>P 43 26 829.3</u>	<u>DE</u>	<u>10/8/93</u>	Priority Claimed
	(Number)	(Country)	(Day/Month/Year Filed)	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
	<u>P 44 18 222.8</u>	<u>DE</u>	<u>25/5/94</u>	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
	(Number)	(Country)	(Day/Month/Year Filed)	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
	<u>P 44 20 157.5</u>	<u>DE</u>	<u>9/6/94</u>	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
	(Number)	(Country)	(Day/Month/Year Filed)	<input type="checkbox"/> Yes <input type="checkbox"/> No
	(Number)	(Country)	(Day/Month/Year Filed)	

(See Note C on back of this page)

☐ See attached list for additional prior foreign applications

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) or PCT International application(s) designating the United States of America listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior application(s) in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, §1.56(a) which occurred between the filing date of the prior application and the national or PCT International filing date of this application:

(List prior U.S. Applications or PCT International applications designating the U.S.)	<u>(Application Serial No.)</u>	<u>(Filing Date)</u>	<u>(Status) (patented, pending, abandoned)</u>
	<u>(Application Serial No.)</u>	<u>(Filing Date)</u>	<u>(Status) (patented, pending, abandoned)</u>

And I hereby appoint as principal attorneys David T. Nikaido, Reg. No. 22,663; Charles M. Marmelstein, Reg. No. 25,895; George E. Oram, Jr., Reg. No. 27,931; Robert B. Murray, Reg. No. 22,980; Martin S. Postman, Reg. No. 18,570; E. Marole Ennas, Reg. No. 32,131; Michael G. Gilman, Reg. No. 19,114; Douglas H. Goldhush, Reg. No. 33,125; Kevin C. Brown, Reg. No. 32,402; Monica Chin Kitts, Reg. No. 36,105; Sharon L. Nolan, Reg. No. 36,335; and John R. Fuisz, Reg. No. 37,327.

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

(See Note D on back of this page)

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29.9.1994
Date

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 Inventor's signature: Richard Lambeth 89.9.1994
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Full name of fourth joint inventor, if any _____
 Inventor's signature _____ Date _____
 Residence _____
 Citizenship _____
 Post Office Address _____

Full name of fifth joint inventor, if any _____
 Inventor's signature _____ Date _____
 Residence _____
 Citizenship _____
 Post Office Address _____

Pull name of sixth joint inventor, if any _____
 Inventor's signature _____ Date _____
 Residence _____
 Citizenship _____
 Post Office Address _____

Full name of seventh joint inventor, if any _____
 Inventor's signature _____ Date _____
 Residence _____
 Citizenship _____
 Post Office Address _____

Full name of eighth joint inventor, if any _____
 Inventor's signature _____ Date _____
 Residence _____
 Citizenship _____
 Post Office Address _____

Full name of ninth joint inventor, if any _____

Inventor's signature _____ Date _____

Residence _____

Citizenship _____

Post Office Address _____

SEQ ID NO.1

TYPE OF SEQUENCE: nucleic acid sequence

NAME AND ORIGIN: MP-52 DNA

LENGTH: 2703 nucleotides

CCATGCGCTCGAAAGGGCAGCGGTGATTTTTTTCACATAAATATATCGCACTTAAATGAG
TTTAGACAGCATGACATCAGAGAGTAATTAATTTGGTTTGGAAATTCGGTTTCCAA
TTCCIGAGTTCAGGTTTGTAAGAAATTTTTCTGAGCACTCGAGGCTGTGAGTGTGTGT
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CGGTTGGCTTTCTCTTTCAAGAACAGGTATTTTTCAGCTGCTGACTGAGAACGGTGCAC
GTCTGTATACGAGAGCATTTCCACTATGGGACTGTGATACAAACACACACCCGCGACACTT
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CCAGGACACCCCTCCCAACAGGCGAGGCTACAGCCCGGACTGTGACCCCAAGAGCAG
CTTCCCGAGGCGAGGCACCCCCAAAAGCAGGATCTGTCCCGAGCTCCTTCTGCTGAAG
AAGGCCAGGAGCCCCCGCCCCCGACAGAGGCCAAGGAGCGGTTTTCGCCCAACCCCAATC
ACACCCCAAGATACATGCTCTCTGCTGACAGGAGCTGTCCGATGCTGACAGAAAGGGA
GGCAACAGCAGCGGTGAAGTTGAGGGCTGGCCCTGGCCACACCATCAACAGCTTTATTGAC
AAAGGCGAAGATGACCGAGTCCCGTGTTCAGGAAGCAGAGGTACGTGTTTGAACATTAGT
GCGCTGAGAGAGGATGGGCTGCTGGGGCGCGAGCTGCGGATCTTTCGGAAGAGCGCTCG
GACACGGCCAGGCCAGCGCCCCCGGAGGCGGGCGGGCTGCCAGCTGAGCTGTCCAGC
TGCCCCAGCGCGCGCGAGCGCGCTCTCTGCTGATGTGCGCTCCGTGCGAGGCTTGAC
GATCTGGCTGGGAGGTGTTCGACATCTGGAAGCTCTTCCGAAACTTTAAGAACTCGGCC
CAGCTGTGCTGAGCTGAGGCGCTGGGAAACGGGGCGAGGCGCTGAGACCTCCGTGGCGCTG
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[illegible]

SEQ ID NO:2

TYPE OF SEQUENCE: amino acid sequence

NAME AND ORIGIN: MP-52 protein

LENGTH: 501 amino acids

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PPLARNVFRP GGHSYGGGAT NANARAKGGT GQTGGLTQPK KDEPKKLPPR
PGGPEPKPGH PPQTRQATAR TVTPKGQLPG GKAPPKAGSV PSSFLLKKAR
EPGPPREPKE PFRPPPITPH EYMLSLYRTL SDADRKGGNS SVKLEAGLAN
TITSFIDKGQ DDRGPVVRKQ RYVFDISALE KDGLLGAE LR ILRKKPSDTA
KPAAPGGGRA AQLKLSSCP S GRQPASLLDV RSVPGLDGSG WEVFDIWKLF
RNFKNSAQLC LELEAWERGR AVDLRGLGFD RAAQVHEKA LFLVFGRTKK
RDLFFNEIKA RSGQDDKTVY EYLFSQRRKR RAPLATRQ GK RPSKNLKARC
SRKALHVNFK DMGWDDWIIA PLEYEAFHCE GLCEFPPLRSH LEPTNHAVIQ
TLMNSMDPES TPPTCCVPT R LSPISILFID SANNVVYKQY EDMVVESCGC R

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT: Hötten, Gertrud
Neidhardt, Helge
Paulista, Michael
- (ii) TITLE OF INVENTION: NEW GROWTH/DIFFERENTIATING FACTOR OF
THE TGF- FAMILY
- (iii) NUMBER OF SEQUENCES: 40
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(C) CITY: Washington
(D) STATE: D.C.
(E) COUNTRY: U.S.A.
(F) ZIP: 20005-5701
- (v) COMPUTER READABLE FORM:
(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1.0, Version #1.25
- (vi) CURRENT APPLICATION DATA:
(A) APPLICATION NUMBER: UNASSIGNED
(B) FILING DATE: 31-AUG-1999
(C) CLASSIFICATION:
- (vii) PRIOR APPLICATION DATA:
(A) APPLICATION NUMBER: 08/288,508
(B) FILING DATE: 10-AUG-1994
- (vii) PRIOR APPLICATION DATA:
(A) APPLICATION NUMBER: DE P 43 26 829.3
(B) FILING DATE: 10-AUG-1993
- (vii) PRIOR APPLICATION DATA:
(A) APPLICATION NUMBER: DE P 44 18 222.8
(B) FILING DATE: 25-MAY-1994
- (vii) PRIOR APPLICATION DATA:
(A) APPLICATION NUMBER: DE P 44 20 157.5
(B) FILING DATE: 09-JUN-1994
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(B) REGISTRATION NUMBER: 36,105
(C) REFERENCE/DOCKET NUMBER: P564-9022
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(B) TELEFAX: (202) 638-4810

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2703 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

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GGGGGAAAAA AAAACTGGAG CACACAGGCA GCATTACGCC ATTCTTCCTT CTGGAAGAAA	300
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GCCCTGACT TGGGCCAGAG ACCCCAGGGG ACCAGGCCAG GATTGGCCAA AGCAGAGGCC	780
AAGGAGAGGC CCCCCTTGGC CCGGAACGTC TTCAGGCCAG GGGGTCACAG CTATGGTGGG	840
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GAGGATTGCT	GCCGCTGTC	TGATGTGACC	AGTGGCAGGC	ACAGGTCCAG	GGAGACAGAC	2400
TCTGAATGGG	ACTGAGTCCC	AGGAAACAGT	GCTTTCGAT	GAGACTCAGC	CCACCAITTC	2460
TCCTCACCTG	GGCCTTCTCA	GCCTCTGGAC	TCTCCTAAGC	ACCTCTCAGG	AGAGCCACAG	2520
GTGCCACTGC	CTCCTCAAAT	CACATTTGTG	CCTGGTGACT	TCCTGTCCCT	GGGACAGTTG	2580
AGAAGCTGAC	TGGGCAAGAG	TGGGAGAGAA	GAGGAGAGGG	CTTGATAGA	GTGAGGAGT	2640
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CCT						2703

(2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 501 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met	Arg	Leu	Pro	Lys	Leu	Leu	Thr	Phe	Leu	Leu	Trp	Tyr	Leu	Ala	Trp
1				5					10					15	

Leu	Asp	Leu	Glu	Phe	Ile	Cys	Thr	Val	Leu	Gly	Ala	Pro	Asp	Leu	Gly
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Glu	Arg	Pro	Pro	Leu	Ala	Arg	Asn	Val	Phe	Arg	Pro	Gly	Gly	His	Ser				
50						55						60							
Tyr	Gly	Gly	Gly	Ala	Thr	Asn	Ala	Asn	Ala	Arg	Ala	Lys	Gly	Gly	Thr				
65			70						75			80							
Gly	Gln	Thr	Gly	Gly	Leu	Thr	Gln	Pro	Lys	Lys	Asp	Glu	Pro	Lys	Lys				
			85						90			95							
Leu	Pro	Pro	Arg	Pro	Gly	Gly	Pro	Glu	Pro	Lys	Pro	Gly	His	Pro	Pro				
			100			105						110							
Gln	Thr	Arg	Gln	Ala	Thr	Ala	Arg	Thr	Val	Thr	Pro	Lys	Gly	Gln	Leu				
115						120						125							
Pro	Gly	Gly	Lys	Ala	Pro	Pro	Lys	Ala	Gly	Ser	Val	Pro	Ser	Ser	Phe				
130						135			140										
Leu	Leu	Lys	Lys	Ala	Arg	Glu	Pro	Gly	Pro	Pro	Arg	Glu	Pro	Lys	Glu				
145			150						155			160							
Pro	Phe	Arg	Pro	Pro	Pro	Ile	Thr	Pro	His	Glu	Tyr	Met	Leu	Ser	Leu				
			165						170			175							
Tyr	Arg	Thr	Leu	Ser	Asp	Ala	Asp	Arg	Lys	Gly	Gly	Asn	Ser	Ser	Val				
180						185						190							
Lys	Leu	Glu	Ala	Gly	Leu	Ala	Asn	Thr	Ile	Thr	Ser	Phe	Ile	Asp	Lys				
195						200						205							
Gly	Gln	Asp	Asp	Arg	Gly	Pro	Val	Val	Arg	Lys	Gln	Arg	Tyr	Val	Phe				
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Asp	Ile	Ser	Ala	Leu	Glu	Lys	Asp	Gly	Leu	Leu	Gly	Ala	Glu	Leu	Arg				
225			230						235			240							
Ile	Leu	Arg	Lys	Lys	Pro	Ser	Asp	Thr	Ala	Lys	Pro	Ala	Ala	Pro	Gly				
			245						250			255							
Gly	Gly	Arg	Ala	Ala	Gln	Leu	Lys	Leu	Ser	Ser	Cys	Pro	Ser	Gly	Arg				
260						265						270							
Gln	Pro	Ala	Ser	Leu	Leu	Asp	Val	Arg	Ser	Val	Pro	Gly	Leu	Asp	Gly				
275						280						285							
Ser	Gly	Trp	Glu	Val	Phe	Asp	Ile	Trp	Lys	Leu	Phe	Arg	Asn	Phe	Lys				
290						295						300							
Asn	Ser	Ala	Gln	Leu	Cys	Leu	Glu	Leu	Glu	Ala	Trp	Glu	Arg	Gly	Arg				
305			310						315			320							
Ala	Val	Asp	Leu	Arg	Gly	Leu	Gly	Phe	Asp	Arg	Ala	Ala	Arg	Gln	Val				
			325						330			335							

His Glu Lys Ala Leu Phe Leu Val Phe Gly Arg Thr Lys Lys Arg Asp
 340 345 350
 Leu Phe Phe Asn Glu Ile Lys Ala Arg Ser Gly Gln Asp Asp Lys Thr
 355 360
 Val Tyr Glu Tyr Leu Phe Ser Gln Arg Arg Lys Arg Arg Ala Pro Leu
 370 375 380
 Ala Thr Arg Gln Gly Lys Arg Pro Ser Lys Asn Leu Lys Ala Arg Cys
 385 390 395 400
 Ser Arg Lys Ala Leu His Val Asn Phe Lys Asp Met Gly Trp Asp Asp
 405 410 415
 Trp Ile Ile Ala Pro Leu Glu Tyr Glu Ala Phe His Cys Glu Gly Leu
 420 425 430
 Cys Glu Phe Pro Leu Arg Ser His Leu Glu Pro Thr Asn His Ala Val
 435 440 445
 Ile Gln Thr Leu Met Asn Ser Met Asp Pro Glu Ser Thr Pro Pro Thr
 450 455 460
 Cys Cys Val Pro Thr Arg Leu Ser Pro Ile Ser Ile Leu Phe Ile Asp
 465 470 475 480
 Ser Ala Asn Asn Val Val Tyr Lys Gln Tyr Glu Asp Met Val Val Glu
 485 490 495
 Ser Cys Gly Cys Arg
 500

(2) INFORMATION FOR SEQ ID NO:3:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 24 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

AGAATTCGCA TGCCATGGTC GACG

24

(2) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 23 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

CTTGAGTACG AGGCTTTCCA CTG

23

(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 24 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

ATTGCGCATGC CATGGTCGAC GAAG

24

(2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 23 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

GGAGCCCACG AATCATGCAG TCA

23

(2) INFORMATION FOR SEQ ID NO:7:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 23 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

ACAGCAGGTG GGTGGTGIGG ACT

23

(2) INFORMATION FOR SEQ ID NO:8:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 44 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

AGAATTGCA TGCCATGTC GACGAAGCTT TTTTTTTTTT TTTT

44

(2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 20 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

CCAGCAGCCC ATCCTTCTCC

20

(2) INFORMATION FOR SEQ ID NO:10:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 24 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

TCCAGGGCAC TAATGTCAAA CACG

24

(2) INFORMATION FOR SEQ ID NO:11:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 24 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

ACTAATGTCA AACACGTACC TCTG

24

(2) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 10 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

AGCGGCCGCT

10

(2) INFORMATION FOR SEQ ID NO:13:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 102 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

Cys Ser Arg Lys Ala Leu His Val Asn Phe Lys Asp Met Gly Trp Asp
1 5 10 15

Asp Trp Ile Ile Ala Pro Leu Glu Tyr Glu Ala Phe His Cys Glu Gly
20 25 30

Leu Cys Glu Phe Pro Leu Arg Ser His Leu Glu Pro Thr Asn His Ala
35 40 45

Val Ile Gln Thr Leu Met Asn Ser Met Asp Pro Glu Ser Thr Pro Pro
50 55 60

Thr Cys Cys Val Pro Thr Arg Leu Ser Pro Ile Ser Ile Leu Phe Ile
65 70 75 80

Asp Ser Ala Asn Asn Val Val Tyr Lys Gln Tyr Glu Asp Met Val Val
85 90 95

Glu Ser Cys Gly Cys Arg
100

(2) INFORMATION FOR SEQ ID NO:14:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 102 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(ix) FEATURE:

- (A) NAME/KEY: misc_feature
- (B) LOCATION: 60
- (C) OTHER INFORMATION: /note= "Xaa is a gap inserted only for comparison purposes"

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

Cys Lys Arg His Pro Leu Tyr Val Asp Phe Ser Asp Val Gly Trp Asn
1 5 10 15

Asp Trp Ile Val Ala Pro Pro Gly Tyr His Ala Phe Tyr Cys His Gly
20 25 30

Glu Cys Pro Phe Pro Leu Ala Asp His Leu Asn Ser Thr Asn His Ala
35 40 45

Ile Val Gln Thr Leu Val Asn Ser Val Asn Ser Xaa Lys Ile Pro Lys
50 55 60

Ala Cys Cys Val Pro Thr Glu Leu Ser Ala Ile Ser Met Leu Tyr Leu
65 70 75 80

Asp Glu Asn Glu Lys Val Val Leu Lys Asn Tyr Gln Asp Met Val Val
85 90 95

Glu Gly Cys Gly Cys Arg
100

(2) INFORMATION FOR SEQ ID NO:15:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 102 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(ix) FEATURE:

- (A) NAME/KEY: misc_feature
- (B) LOCATION: 60
- (C) OTHER INFORMATION: /note= "Xaa is a gap inserted only for comparison purposes"

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

Cys Arg Arg His Ser Leu Tyr Val Asp Phe Ser Asp Val Gly Trp Asn
 1 5 10 15
 Asp Trp Ile Val Ala Pro Pro Gly Tyr Gln Ala Phe Tyr Cys His Gly
 20 25 30
 Asp Cys Pro Phe Pro Leu Ala Asp His Leu Asn Ser Thr Asn His Ala
 35 40 45
 Ile Val Gln Thr Leu Val Asn Ser Val Asn Ser Xaa Ser Ile Pro Lys
 50 55 60
 Ala Cys Cys Val Pro Thr Glu Leu Ser Ala Ile Ser Met Leu Tyr Leu
 65 70 75 80
 Asp Glu Tyr Asp Lys Val Val Leu Lys Asn Tyr Gln Glu Met Val Val
 85 90 95
 Glu Gly Cys Gly Cys Arg
 100

(2) INFORMATION FOR SEQ ID NO:16:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 102 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

Cys Lys Lys His Glu Leu Tyr Val Ser Phe Arg Asp Leu Gly Trp Gln
 1 5 10 15
 Asp Trp Ile Ile Ala Pro Glu Gly Tyr Ala Ala Phe Tyr Cys Asp Gly
 20 25 30
 Glu Cys Ser Phe Pro Leu Asn Ala His Met Asn Ala Thr Asn His Ala
 35 40 45
 Ile Val Gln Thr Leu Val His Leu Met Phe Pro Asp His Val Pro Lys
 50 55 60
 Pro Cys Cys Ala Pro Thr Lys Leu Asn Ala Ile Ser Val Leu Tyr Phe
 65 70 75 80
 Asp Asp Ser Ser Asn Val Ile Leu Lys Lys Tyr Arg Asn Met Val Val
 85 90 95
 Arg Ser Cys Gly Cys His
 100

(2) INFORMATION FOR SEQ ID NO:17:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 102 amino acids

(B) TYPE: amino acid
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

```

Cys Arg Lys His Glu Leu Tyr Val Ser Phe Gln Asp Leu Gly Trp Gln
 1           5           10           15
Asp Trp Ile Ile Ala Pro Lys Gly Tyr Ala Ala Asn Tyr Cys Asp Gly
 20           25           30
Glu Cys Ser Phe Pro Leu Asn Ala His Met Asn Ala Thr Asn His Ala
 35           40           45
Ile Val Gln Thr Leu Val His Leu Met Asn Pro Glu Tyr Val Pro Lys
 50           55           60
Pro Cys Cys Ala Pro Thr Lys Leu Asn Ala Ile Ser Val Leu Tyr Phe
 65           70           75           80
Asp Asp Asn Ser Asn Val Ile Leu Lys Lys Tyr Arg Asn Met Val Val
 85           90           95
Arg Ala Cys Gly Cys His
100

```

(2) INFORMATION FOR SEQ ID NO:18:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 102 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

```

Cys Lys Lys His Glu Leu Tyr Val Ser Phe Arg Asp Leu Gly Trp Gln
 1           5           10           15
Asp Trp Ile Ile Ala Pro Glu Gly Tyr Ala Ala Tyr Tyr Cys Glu Gly
 20           25           30
Glu Cys Ala Phe Pro Leu Asn Ser Tyr Met Asn Ala Thr Asn His Ala
 35           40           45
Ile Val Gln Thr Leu Val His Phe Ile Asn Pro Glu Thr Val Pro Lys
 50           55           60
Pro Cys Cys Ala Pro Thr Gln Leu Asn Ala Ile Ser Val Leu Tyr Phe
 65           70           75           80
Asp Asp Ser Ser Asn Val Ile Leu Lys Lys Tyr Arg Asn Met Val Val
 85           90           95
Arg Ala Cys Gly Cys His
100

```

(2) INFORMATION FOR SEQ ID NO:19:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 36 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

ATGAATTCCTCC ATGGACCTGG GCTGGMAGKA MTGGAT

36

(2) INFORMATION FOR SEQ ID NO:21:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

ACGTGGGGTG GAATGACTGG AT

22

(2) INFORMATION FOR SEQ ID NO:21:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

ATATTGGCTG GAGTGAATGG AT

22

(2) INFORMATION FOR SEQ ID NO:22:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

ATGTTGGCTG GAATGACTGG AT

22

(2) INFORMATION FOR SEQ ID NO:23:

- (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 22 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

ACCTGGGCTG GCAGGACTGG AT

22

(2) INFORMATION FOR SEQ ID NO:24:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

AGGACCTCGG CTGGAAGTGG AT

22

(2) INFORMATION FOR SEQ ID NO:25:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

GGGATCTAGG GTGGAAATGG AT

22

(2) INFORMATION FOR SEQ ID NO:26:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

AGGATCTGGG CTGGAAGTGG GT

22

(2) INFORMATION FOR SEQ ID NO:27:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

AGCTGGGCTG GGAACGGTGG AT

22

(2) INFORMATION FOR SEQ ID NO:28:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

ACATCGGCTG GAATGACTGG AT

22

(2) INFORMATION FOR SEQ ID NO:29:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

TCATCGGCTG GAACTGCTGG AT

22

(2) INFORMATION FOR SEQ ID NO:30:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 29 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

ATGAATTCGA GCTGCGTSGG SRCACAGCA

29

(2) INFORMATION FOR SEQ ID NO:31:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 21 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

GAGTTCGTGC GGGACACAGC A

21

(2) INFORMATION FOR SEQ ID NO:32:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 21 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:

CATCTTTTCT GGTACACAGC A

21

(2) INFORMATION FOR SEQ ID NO:33:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 21 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

CAGTTCAGTG GGCACACAAC A

21

(2) INFORMATION FOR SEQ ID NO:34:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 21 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:

GAGCTGCGTG GCGCACAGC A

21

(2) INFORMATION FOR SEQ ID NO:35:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 21 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

CAGCGCCTGC GGCACGCAGC A

21

(2) INFORMATION FOR SEQ ID NO:36:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 21 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

TAAATCTTGG GACACGCAGC A

21

(2) INFORMATION FOR SEQ ID NO:37:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 21 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

CAGGTCCTGG GGCACGCAGC A

21

(2) INFORMATION FOR SEQ ID NO:38:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 21 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:

CCCTGGGAGA GCAGCACAGC A

21

(2) INFORMATION FOR SEQ ID NO:39:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 21 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

CAGCTTGGTG GGCACACAGC A

21

(2) INFORMATION FOR SEQ ID NO:40:

- (i) SEQUENCE CHARACTERISTICS:

